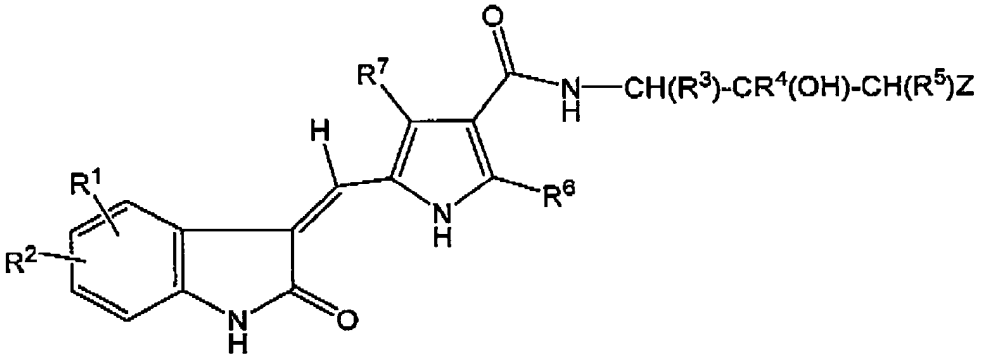
Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

1 – 17. (Canceled)

18. (Currently Amended) A method of synthesizing a compound of Formula (I):



(I)

wherein:

R^1 is selected from the group consisting of hydrogen, halo, alkyl, haloalkoxy, ~~cycloalkyl~~ cycloalkyl, aryl, heteroaryl, heteroalicyclic, hydroxy, alkoxy, -$(CO)R^8$, -NR^9R^{10}, -$(CHR^3)_nR^{11}$ and -$C(O)NR^{12}R^{13}$;

R^2 is selected from the group consisting of hydrogen, halo, alkyl, trihalomethyl, hydroxy, alkoxy, cyano, -NR^9R^{10}, -$NR^9C(O)R^{10}$, -$C(O)R^8$, aryl, heteroaryl, –$S(O)_2NR^9R^{10}$ and -SO_2R^{14} ~~(wherein R^{14} is alkyl, aryl, aralkyl, heteroaryl and heteroaralkyl)~~;

R^3, R^4 and R^5 are independently hydrogen or alkyl;

Z is aryl, heteroaryl, heterocycle, or –$NR^{15}R^{16}$ wherein R^{15} and R^{16} are independently hydrogen or alkyl; or R^{15} and R^{16} together with the nitrogen atom to which they are attached ~~from~~ form a heterocycloamino group;

R^6 is selected from the group consisting of hydrogen ~~or~~ and alkyl;

R^7 is selected from the group consisting of hydrogen, alkyl, aryl, heteroaryl, and -$C(O)R^{17}$ as defined below;

R^8 is selected from the group consisting of hydrogen, hydroxy, alkoxy and aryloxy;

R^9 and R^{10} are independently selected from the group consisting of hydrogen, alkyl, cyanoalkyl, cycloalkyl, aryl and heteroaryl; or

R^9 and R^{10} combine to form a heterocycloamino group;

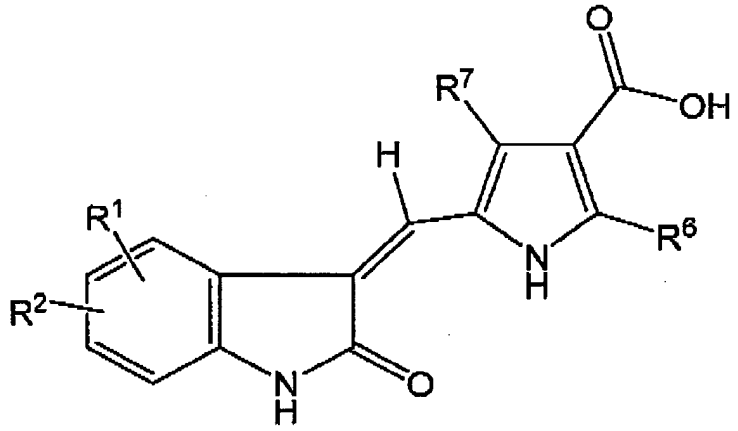
R^{11} is selected from the group consisting of hydroxy, -$C(O)R^8$, -NR^9R^{10} and -$C(O)NR^9R^{10}$ wherein R^8, R^9 and R^{10} are as defined above;

R^{12} and R^{13} are independently selected from the group consisting of hydrogen, alkyl, hydroxyalkyl, and aryl; or R^{12} and R^{13} together with the nitrogen atom to which they are attached form a heterocycloamino; and

R^{17} is selected from the group consisting of alkyl, cycloalkyl, aryl and heteroaryl

comprising reacting

a compound of Formula (II)



(II)

with

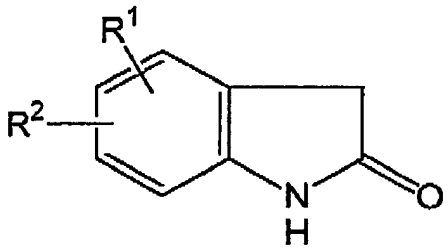
a compound of Formula (III)

(III)

in the presence of an organic solvent and a coupling agent, to form compound (I), wherein R^1, R^2, R^3, R^4, R^5, R^7 and Z are as defined above.

19. (Currently Amended) The method of claim 18, wherein compound (II) is formed by reacting

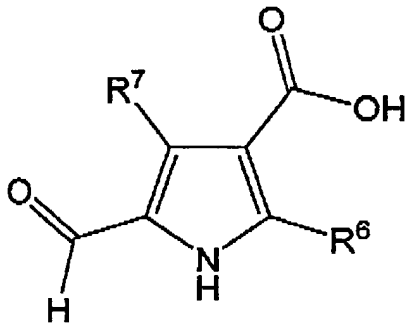
a compound of Formula (IV)



(IV)

with

a compound of Formula (V)



(V)

in the presence of a solvent and a base, wherein, R^1, R^2, R^6 and R^7 are as defined above.

20. (Canceled)

21. (Currently Amended) The method of claim ~~20~~ 18, wherein the organic solvent is ~~dimethylforamide~~ dimethylformamide or tetrahydrofuran.

22. (Currently Amended) The method of claim ~~20~~ 18, wherein the coupling agent is dicyclohexylcarbodiimide, DEAD, EDC or HOBt.

23. (Canceled)

24. (Previously Presented) A method of synthesizing 5-[5-fluoro-2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-morpholin-4-yl-propyl)-amide comprising:

reacting morpholino and epichlorohydrin to form 1-chloro-3-morpholin-4-yl-propan-2-ol;

reacting 1-chloro-3-morpholin-4-yl-propan-2-ol with ammonia to form 1-amino-3-morpholin-4-yl-propan-2-ol;

reacting 1-amino-3-morpholin-4-yl-propan-2-ol with 5-(5-Fluoro-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid to form 5-[5-fluoro-2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-morpholin-4-yl-propyl)-amide.

25 - 28. (Canceled)

29. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
5-[5-fluoro-2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-morpholin-4-yl-propyl)-amide for Formula (I),
5-(5-Fluoro-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3-morpholin-4-yl-propan-2-ol for Formula (III).

30. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
2,4-dimethyl-5-[2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-morpholin-4-yl-propyl)-amide for Formula (I),
5-(2-Oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3-morpholin-4-yl-propan-2-ol for Formula (III).

31. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
5-[5-chloro-2-oxo-1,2-dihydro-indol-(3Z)-ylidene-methyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-morpholin-4-yl-propyl)-amide for Formula (I),
5-(5-Chloro-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3-morpholin-4-yl-propan-2-ol for Formula (III).

32. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
5-[5-bromo-2-oxo-1,2-dihydro-indol-(3Z)-ylidene-methyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-morpholin-4-yl-propyl)-amide for Formula (I),
5-(5-Bromo-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3-morpholin-4-yl-propan-2-ol for Formula (III).

33. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are

2,4-dimethyl-5-[2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-[1,2,3]triazol-1-yl-propyl)-amide for Formula (I),
5-(2-Oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3(1,2,3)triazole-1-yl-propan-2-ol for Formula (III).

34. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
5-[5-fluoro-2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-[1,2,3]triazol-1-yl-propyl)-amide for Formula (I),
5-(5-Fluoro-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3(1,2,3)triazol-1-yl-propan-2-ol for Formula (III).

35. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
5-[5-Chloro-2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-[1,2,3]triazol-1-yl-propyl)-amide for Formula (I),
5-(5-Chloro-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3(1,2,3)triazole-1-yl-propan-2-ol for Formula (III).

36. (Previously Presented) The method of claim 18, wherein compounds of Formulas (I), (II) and (III) are
5-[5-bromo-2-oxo-1,2-dihydro-indol-(3Z)-ylidenemethyl]-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid (2-hydroxy-3-[1,2,3]triazol-1-yl-propyl)-amide for Formula (I),
5-(5-Bromo-2-oxo-1,2-dihydro-indol-3-ylidenemethyl)-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid for Formula (II) and
1-amino-3(1,2,3)triazole-1-yl-propan-2-ol for Formula (III).